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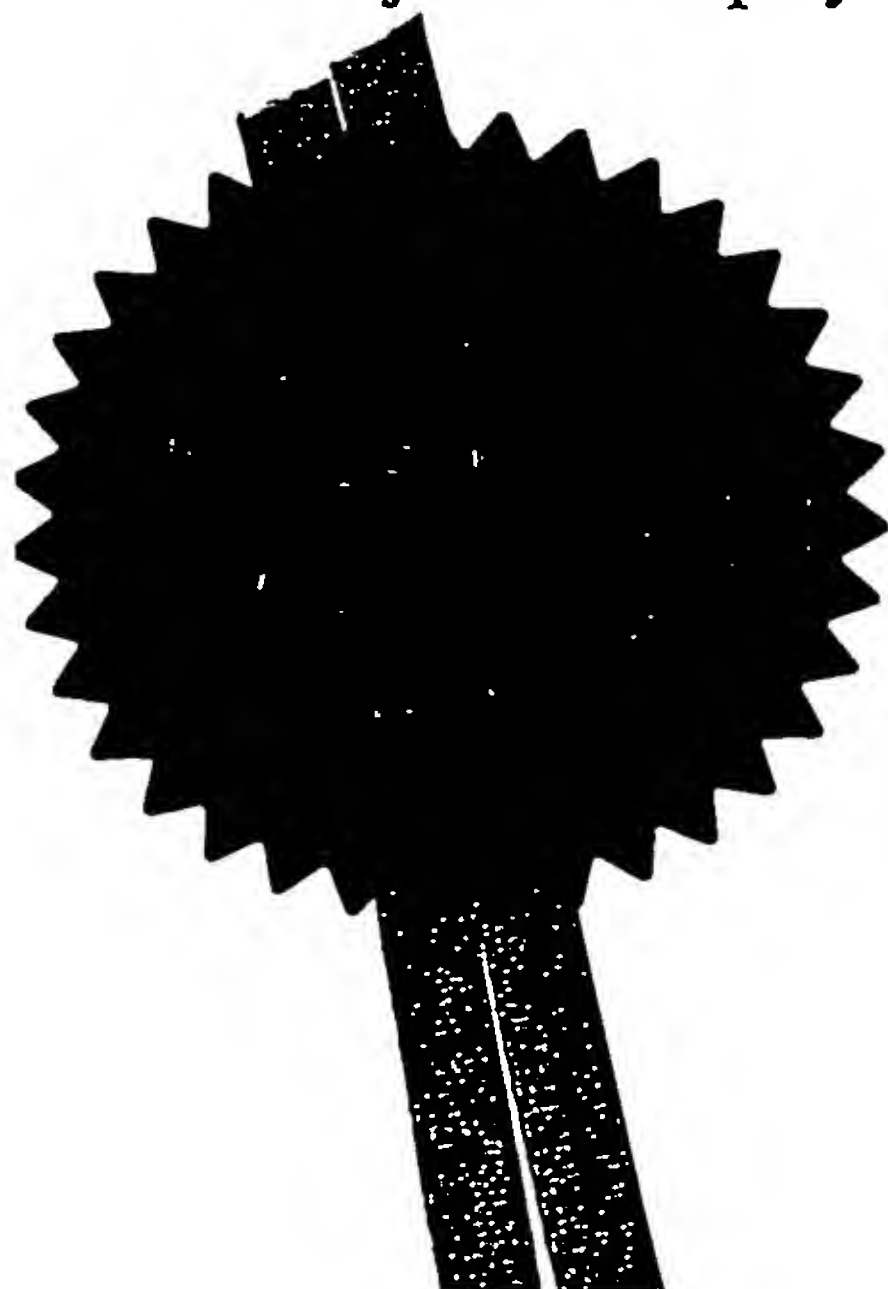
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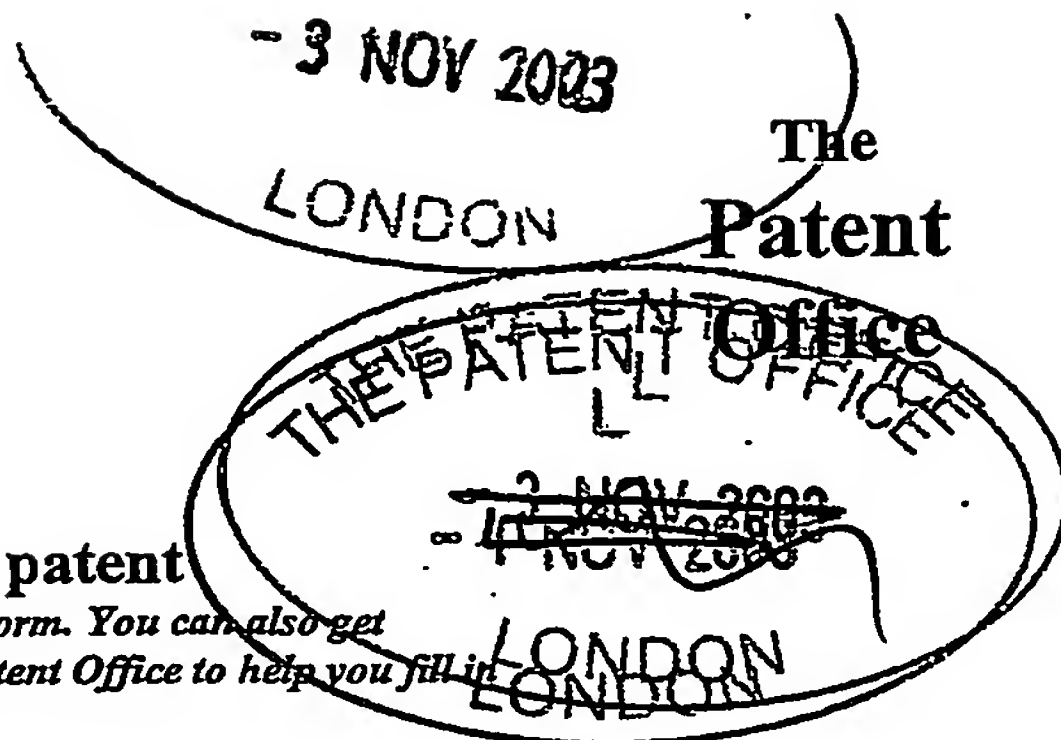
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Patents Form 1/77

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P01/7700 0.00-0325617.9

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1. Your reference C4334 (C)/rkk

2. Patent application number  
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0 3 NOV 2003

0325617.9

3. Full name, address and postcode of the or of each applicant (underline all surnames)

UNILEVER PLC  
UNILEVER HOUSE, BLACKFRIARS  
LONDON, EC4P 4BQ

Patents ADP number (if you know it)

50426956002

If the applicant is a corporate body, give the country/state of its incorporation

UNITED KINGDOM

4. Title of the invention

RED BLEACHING COMPOSITIONS

5. Name of your agent (if you have one)

ELLIOTT, Peter William

"Address for Service" in the United Kingdom to which all correspondence should be sent (including the postcode)

PATENT DEPARTMENT, UNILEVER PLC  
COLWORTH HOUSE, SHARNBROOK  
BEDFORD, MK44 1LQ

Patents ADP number (if you know it)

1628003

6. Priority: Complete this section if you are declaring priority from one or more earlier patent applications, filed in the last 12 months.

Country	Priority application number (if you know it)	Date of filing (day / month / year)
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7. Divisionals, etc: Complete this section only if this application is a divisional application or resulted from an entitlement dispute (see note f)

Number of earlier application	Date of filing (day / month / year)
-------------------------------	-------------------------------------

8. Is a Patents Form 7/77 (Statement of inventorship and of right to grant of a patent) required in support of this request?

Answer YES if:

- a) any applicant named in part 3 is not an inventor, or
  - b) there is an inventor who is not named as an applicant, or
  - c) any named applicant is a corporate body.
- Otherwise answer NO (See note d)

YES

# Patents Form 1/77

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Continuation sheets of this form

Description	24
Claim(s)	2 DC
Abstract	1
Drawing(s)	

10. If you are also filing any of the following, state how many against each item.

## Priority Documents

Translations of priority documents

Statement of inventorship and right to grant of a patent (Patents Form 7/77)

Request for preliminary examination and search (Patents Form 9/77) 1 ✓

Request for substantive examination (Patents Form 10/77)

Any other documents (please specify)

11. I/We request the grant of a patent on the basis of this application.

Signature(s)



Date: 03-11-03

Sandra Jane EDWARDS, Authorised Signatory

12. Name, daytime telephone number and e-mail address, if any, of person to contact in the United Kingdom

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DUPLICATE

- 1 -

## RED BLEACHING COMPOSITIONS

FIELD OF INVENTION

The present invention relates to the photo-bleaching of  
5 products, in particular laundry bleaching.

BACKGROUND OF INVENTION

In a conventional bleaching treatment, a substrate such as a  
laundry fabric or other textile is subjected to hydrogen  
10 peroxide, or to substances which can generate hydroperoxyl  
radicals, such as inorganic or organic peroxides.

A preferred approach to generating hydroperoxyl bleach  
radicals is the use of inorganic peroxides coupled with  
15 organic precursor compounds. These systems are employed for  
many commercial laundry powders. For example, various  
European systems are based on tetraacetyl ethylenediamine  
(TAED) as the organic precursor coupled with sodium  
perborate or sodium percarbonate, whereas in the United  
20 States laundry bleach products are typically based on sodium  
nonanoyloxybenzenesulphonate (SNOBS) as the organic  
precursor coupled with sodium perborate. Alternatively, or  
additionally, hydrogen peroxide and peroxy systems can be  
activated by bleach catalysts, such as by complexes of iron  
25 and the ligand N4Py (i.e. N, N-bis(pyridin-2-yl-methyl)-  
bis(pyridin-2-yl)methylamine) disclosed in WO95/34628.

Another approach to bleaching is that of using radical  
photo-initiators, which are generally organic chemicals  
30 which on absorption of light, particularly UV light, form  
reactive radicals. They are widely used to initiate the

polymerisation of alkenes and thereby cure coatings. They may also be used as photobleach agents as discussed in GB 9917451. Radical photoinitiators, in general, are discussed by H.F. Gruber in *Prog. Polym. Sci.* Vol 17. 953-1044.

5

#### SUMMARY OF INVENTION

We have found that Food red 14 is surprisingly active in photo-bleaching catechol-type stains, polyphenolics-type stains and polycyclic hydroxylated aromatic-type stains.

10 The Food red 14 may be used on its own as the bleaching species or in conjunction with other bleaching species. It is preferred that those other bleaching species, if used, possess different bleaching profiles. The stain bleaching profile of known photo-bleaches is generally that of  
15 ceretenoid type stains, for example tomato stains which is similar to the bleaching profile of "air" bleaching catalysts.

These catechol-type stains, polyphenolics-type stains and  
20 polycyclic hydroxylated aromatic-type stains are as a result of the chromophores found in tea, coffee, blackberry, blueberry, blackcurrant, red wine, banana and the like. These stains are characteristic and distinct from oily food type stains such as tomato oil stain, curry oil stain, mango  
25 stain, annatto derived stain, colorau derived stain, and sebum derived stain etc.

The economic factors in cleaning products are a dominant in many decisions to launch a product. By substituting or  
30 reducing the amounts of more expensive bleaching components with Food red 14 an efficacious product may be produced with



at less cost. This exemplified by the following, Food red 14 exhibits similar bleaching efficacy to that of perborate/percarbonate when used in conjunction with TAED on certain stains. Another benefit is that Food red 14 may be used in washes of lower temperature than that of the peroxy systems. In addition, the level of Food Red 14 exhibits bleaching effects at surprisingly low levels. Of the red dyes Food Red 14 deposited effectively on the textile.

10 The present invention provides a bleaching composition comprising: a) from 0.0001 wt/wt% to 0.1 wt/wt% of Food red 14 dye; b) from 0 to 40% other bleaching species; and, c) the balance carriers and adjunct ingredients to 100 wt/wt % of the total bleaching composition, wherein upon addition of  
15 a unit dose of the bleaching composition to an aqueous environment a wash liquor having a pH in the range 8 to 11 is provided.

The fact that Food Red 14 is brightly red makes its  
20 application in textile bleaching unusual because one would expect their use to result in discoloured rather than whitened textiles. When Food red 14 is used solely as the bleaching species the consumer is reassured in the laundry context that the bleach used is mild because of that fact  
25 that it is an acceptable a food additive. Food dyes are used to enhance the visual appearance of many foods. As humans consume them, they undergo rigorous testing and examination to ensure they represent no risk to health. In addition, a solution of Food Red 14 under indoor lighting is much paler  
30 to the eye than that of Rose Bengal, which has an intense colour in solution, despite having a lower optical density

than Food red 14. This permits higher dosages of Food red 14 to be used than, for example, Rose Bengal without overly colouring the liquor.

- 5 The present invention also extends to a method of photo-bleaching a textile.

The light used for photo-bleaching may be sunlight, florescent light or that from an ordinary light bulb.

10

Photo-bleaching is still effective where the concentration of Food red 14 on the textile/cloth is so low as to be barely perceptible to the human eye. The Food red 14 if perceptible rapidly photo-fades.

15

The present invention may be applied locally, suitable pre-treatment means for application include sprays, pens, roller-ball devices, bars, soft solid applicator sticks.

- 20 A unit dose as used herein is a particular amount of the bleaching composition used for a type of wash. The unit dose may be in the form of a defined volume of powder, granules or tablet.

25 DETAILED DESCRIPTION OF THE INVENTION

Food red 14 (Erythrosine B) (E 127; CI number 45430) (CAS 16423-68-0) is a widely used red food colouring, see Hunger K. Industrial Dyes: Chemistry Properties And Applications. Wiley-VCH, Heidelberg 2003.

30

The concentration of Food red 14 in a wash liquor is from 5ppb to 1000ppm, preferably 10ppb to 100ppm, more preferably 50ppb to 5ppm, and most preferably 100ppb to 1ppm. A same concentration may also be used for a spot treatment of a stain or of a commercial liquid formulation. A suitable concentration in a powder detergent would be 0.0001 wt/wt% to 0.1 wt/wt%, most preferred is 0.001 wt/wt% to 0.01 wt/wt%.

#### 10 Other Bleaching Species

The bleaching composition may also contain other bleaching components, for example other photo-bleaches, a transition metal catalyst which is present in a bleaching composition that is substantially devoid of peroxy species, and peroxy bleaching systems. An example of a preferred additional photo-bleach is vitamin K3.

The use of bleaching catalysts for stain removal has been developed over recent years and may be used in the present invention. Examples of transition metal catalysts that may be used are found, for example, in: WO0060045, WO0248310, WO0029537 and WO0012667. The catalysts may be used for catalysing peroxy or "air" bleaching as described in WO0248301. The catalyst may alternatively be provided as the free ligand that forms a complex in situ.

The bleaching composition when use as an "air" bleaching composition is substantially devoid of a peroxygen bleach or a peroxy-based or peroxy-generating bleach system. It is believed that adventitious hydroperoxides within an oily stain serve to bleach the stain together with the catalyst.



The term "substantially devoid of a peroxygen bleach or a peroxy-based or peroxy-generating bleach system" should be construed within spirit of the invention. It is preferred that the composition has as low a content of peroxy species present as possible. It is preferred that the bleaching formulation contains less than 1 % wt/wt total concentration of peracid or hydrogen peroxide or source thereof, preferably the bleaching formulation contains less than 0.3 % wt/wt total concentration of peracid or hydrogen peroxide or source thereof, most preferably the bleaching composition is devoid of peracid or hydrogen peroxide or source thereof.

Conversely the bleaching composition may be used as a peroxy bleaching composition where the bleaching composition comprises a peroxy bleaching species. The peroxy bleaching species may be a compound, which is capable of yielding hydrogen peroxide in aqueous solution. Hydrogen peroxide sources are well known in the art. They include the alkali metal peroxides, organic peroxides such as urea peroxide, and inorganic persalts, such as the alkali metal perborates, percarbonates, perphosphates persilicates and persulphates. Mixtures of two or more such compounds may also be suitable.

Particularly preferred are sodium perborate tetrahydrate and, especially, sodium perborate monohydrate. Sodium perborate monohydrate is preferred because of its high active oxygen content. Sodium percarbonate may also be preferred for environmental reasons. The amount thereof in the composition of the invention usually will be within the range of about 1-35% by weight, preferably from 5-25% by

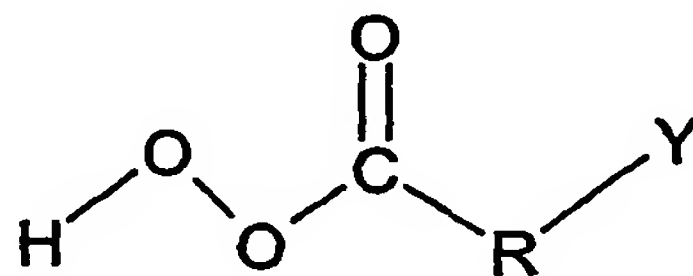
weight. One skilled in the art will appreciate that these amounts may be reduced in the presence of a bleach precursor e.g., N,N,N',N'-tetraacetyl ethylene diamine (TAED).

- 5 Another suitable hydrogen peroxide generating system is a combination of a C1-C4 alkanol oxidase and a C1-C4 alkanol, especially a combination of methanol oxidase (MOX) and ethanol. Such combinations are disclosed in International Application PCT/EP 94/03003 (Unilever), which is  
10 incorporated herein by reference.

Alkylhydroxy peroxides are another class of peroxy bleaching compounds. Examples of these materials include cumene hydroperoxide and t-butyl hydroperoxide.

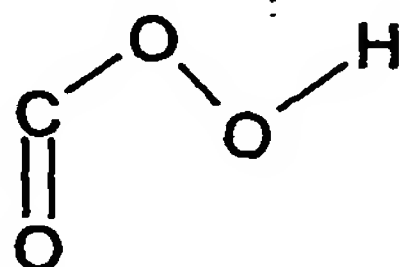
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Organic peroxyacids may also be suitable as the peroxy bleaching compound. Such materials normally have the general formula:



20

wherein R is an alkylene or substituted alkylene group containing from 1 to about 20 carbon atoms, optionally having an internal amide linkage; or a phenylene or substituted phenylene group; and Y is hydrogen, halogen,  
25 alkyl, aryl, an imido-aromatic or non-aromatic group, a COOH or



group or a quaternary ammonium group.

Typical monoperoxy acids useful herein include, for example:

- 5 (i) peroxybenzoic acid and ring-substituted peroxybenzoic acids, e.g. peroxy-alpha-naphthoic acid;
- (ii) aliphatic, substituted aliphatic and arylalkyl monoperoxyacids, e.g., peroxyauric acid, peroxysearic acid  
10 and N,N-phthaloylaminoperoxy caproic acid (PAP); and
- (iii) 6-octylamino-6-oxo-peroxyhexanoic acid.

Typical diperoxyacids useful herein include, for example:

- 15 (iv) 1,12-diperoxydodecanedioic acid (DPDA);
- (v) 1,9-diperoxyazelaic acid;
- 20 (vi) diperoxybrassicic acid; diperoxysebasic acid and diperoxyisophthalic acid;
- (vii) 2-decyldiperoxybutane-1,4-dioic acid; and
- 25 (viii) 4,4'-sulphonylbisperoxybenzoic acid.

Also inorganic peroxyacid compounds are suitable, such as for example potassium monopersulphate (MPS). If organic or inorganic peroxyacids are used as the peroxygen compound,  
30 the amount thereof will normally be within the range of about 2-10% by weight, preferably from 4-8% by weight.

Peroxyacid bleach precursors are known and amply described in literature, such as in the British Patents 836988; 864,798; 907,356; 1,003,310 and 1,519,351; German Patent 3,337,921; EP-A-0185522; EP-A-0174132; EP-A-0120591; and  
5 U.S. Pat. Nos. 1,246,339; 3,332,882; 4,128,494; 4,412,934 and 4,675,393.

Another useful class of peroxyacid bleach precursors is that of the cationic i.e. quaternary ammonium substituted  
10 peroxyacid precursors as disclosed in US Pat. Nos. 4,751,015 and 4,397,757, in EP-A0284292 and EP-A-331,229. Examples of peroxyacid bleach precursors of this class are:

2-(N,N,N-trimethyl ammonium) ethyl sodium-4-sulphonphenyl  
15 carbonate chloride (SPCC);

N-octyl-N,N-dimethyl-N10-carbophenoxy decyl ammonium  
chloride (ODC);

20 3-(N,N,N-trimethyl ammonium) propyl sodium-4-sulphophenyl  
carboxylate; and

N,N,N-trimethyl ammonium toluyloxy benzene sulphonate.

25 A further special class of bleach precursors is formed by the cationic nitriles as disclosed in EP-A-303,520 and in European Patent Specification No.'s 458,396 and 464,880.

Any one of these peroxyacid bleach precursors can be used in  
30 the present invention, though some may be more preferred than others.

Of the above classes of bleach precursors, the preferred classes are the esters, including acyl phenol sulphonates and acyl alkyl phenol sulphonates; the acyl-amides; and the quaternary ammonium substituted peroxyacid precursors  
5 including the cationic nitriles.

Examples of said preferred peroxyacid bleach precursors or activators are sodium-4-benzoyloxy benzene sulphonate (SBOBS); N,N,N',N'-tetraacetyl ethylene diamine (TAED);  
10 sodium-1-methyl-2-benzoyloxy benzene-4-sulphonate; sodium-4-methyl-3-benzoyloxy benzoate; SPCC; trimethyl ammonium toluoyloxy-benzene sulphonate; sodium nonanoyloxybenzene sulphonate (SNOBS); sodium 3,5,5-trimethyl hexanoyl-oxybenzene sulphonate (STHOBS); and the substituted cationic  
15 nitriles.

Other classes of bleach precursors for use with the present invention are found in WO0015750, for example 6-(nonanamidocaproyl)oxybenzene sulphonate.

20

The precursors may be used in an amount of up to 12%, preferably from 2-10% by weight, of the composition.

#### Balance Carriers and Adjunct Ingredients

25

The following provides examples of suitable balance carriers and adjunct ingredients.

These are generally surfactants, builders, foam agents,  
30 anti-foam agents, solvents, and enzymes. The use and amounts of these components are such that the bleaching



composition performs depending upon economics, environmental factors and use of the bleaching composition. It is preferred that the bleaching composition comprises from 5 to 40 wt/wt % of a surfactant having an HLB greater than 15.

5

To that extent, the composition comprises a surfactant and optionally other conventional detergent ingredients. The invention in its second aspect provides an enzymatic detergent composition which comprises from 0.1 - 50 % by weight, based on the total detergent composition, of one or more surfactants. This surfactant system may in turn comprise 0 - 95 % by weight of one or more anionic surfactants and 5 to 100 % by weight of one or more nonionic surfactants. The surfactant system may additionally contain amphoteric or zwitterionic detergent compounds, but this is not normally desired owing to their relatively high cost. The enzymatic detergent composition according to the invention will generally be used as a dilution in water of about 0.05 to 2%.

20

In general, the nonionic and anionic surfactants of the surfactant system may be chosen from the surfactants described "Surface Active Agents" Vol. 1, by Schwartz & Perry, Interscience 1949, Vol. 2 by Schwartz, Perry & Berch, Interscience 1958, in the current edition of "McCutcheon's Emulsifiers and Detergents" published by Manufacturing Confectioners Company or in "Tenside-Taschenbuch", H. Stache, 2nd Edn., Carl Hauser Verlag, 1981.

30 Suitable nonionic detergent compounds which may be used include, in particular, the reaction products of compounds

having a hydrophobic group and a reactive hydrogen atom, for example, aliphatic alcohols, acids, amides or alkyl phenols with alkylene oxides, especially ethylene oxide either alone or with propylene oxide. Specific nonionic detergent

5 compounds are C<sub>6</sub>-C<sub>22</sub> alkyl phenol-ethylene oxide condensates, generally 5 to 25 EO, i.e. 5 to 25 units of ethylene oxide per molecule, and the condensation products of aliphatic C<sub>8</sub>-C<sub>18</sub> primary or secondary linear or branched alcohols with ethylene oxide, generally 5 to 40 EO.

10

Suitable anionic detergent compounds which may be used are usually water-soluble alkali metal salts of organic sulphates and sulphonates having alkyl radicals containing from about 8 to about 22 carbon atoms, the term alkyl being

15

used to include the alkyl portion of higher acyl radicals. Examples of suitable synthetic anionic detergent compounds are sodium and potassium alkyl sulphates, especially those obtained by sulphating higher C<sub>8</sub>-C<sub>18</sub> alcohols, produced for example from tallow or coconut oil, sodium and potassium

20

alkyl C<sub>9</sub>-C<sub>20</sub> benzene sulphonates, particularly sodium linear secondary alkyl C<sub>10</sub>-C<sub>15</sub> benzene sulphonates; and sodium alkyl glyceryl ether sulphates, especially those ethers of the higher alcohols derived from tallow or coconut oil and synthetic alcohols derived from petroleum. The preferred

25

anionic detergent compounds are sodium C<sub>11</sub>-C<sub>15</sub> alkyl benzene sulphonates and sodium C<sub>12</sub>-C<sub>18</sub> alkyl sulphates. Also applicable are surfactants such as those described in

EP-A-328 177 (Unilever), which show resistance to salting-out, the alkyl polyglycoside surfactants described

30

in EP-A-070 074, and alkyl monoglycosides.

Preferred surfactant systems are mixtures of anionic with nonionic detergent active materials, in particular the groups and examples of anionic and nonionic surfactants pointed out in EP-A-346 995 (Unilever). Especially preferred  
5 is surfactant system that is a mixture of an alkali metal salt of a C<sub>16</sub>-C<sub>18</sub> primary alcohol sulphate together with a C<sub>12</sub>-C<sub>15</sub> primary alcohol 3-7 EO ethoxylate.

10 The nonionic detergent is preferably present in amounts greater than 10%, e.g. 25-90% by weight of the surfactant system. Anionic surfactants can be present for example in amounts in the range from about 5% to about 40% by weight of the surfactant system.

15 The composition may contain additional enzymes as found in WO 01/00768 A1 page 15, line 25 to page 19, line 29, the contents of which are herein incorporated by reference.

20 Builders, polymers and other enzymes as optional ingredients may also be present as found in WO0060045.

Suitable detergency builders as optional ingredients may also be present as found in WO0034427.

25 The composition of the present invention may be used for laundry cleaning, hard surface cleaning (including cleaning of lavatories, kitchen work surfaces, floors, mechanical  
30 treatment, pulp bleaching during the manufacture of paper, leather manufacture, dye transfer inhibition, food

processing, starch bleaching, sterilisation, whitening in oral hygiene preparations and/or contact lens disinfection.

5 In the context of the present invention, bleaching should be understood as relating generally to the decolourisation of stains or of other materials attached to or associated with a substrate. However, it is envisaged that the present invention can be applied where a requirement is the removal and/or neutralisation by an oxidative bleaching reaction of  
10 malodours or other undesirable components attached to or otherwise associated with a substrate.

### Examples

#### 15 Example 1

A 1.5g/L stock solution of a base washing powder was created. The washing powder contained 18% NaLAS, 73% salts (silicate, sodium tripolyphosphate, sulphate, carbonate), 3%  
20 minors including perborate, fluoereser and enzymes, remainder impurities and water. The solution was divided into 4, 200ml aliquots. To each was added various amounts of the food red 14, and then 2 BC1 test clothes and 4 white cotton clothes. All clothes were the same size and weight, the total weight  
25 of cloth added was 5g. BC1 is a standard stained test cloth used in laundry evaluation and is a model tea stain monitor. BC1 clothes may be purchased from e.g., the center for test materials, Vlaardingen, NL.

30 The clothes were then agitated in solution for 20 mins, removed rinsed and the 2 BC1 clothes and 2 of the white

clothes irradiated in a weatherometer (WOM) for 30 minutes. A WOM produces artificial sunlight and was set up to give  $385 \text{ W/m}^2$  in the UV-visible range. The remaining 2 white clothes were dried in the dark in a tumble drier.

5

Following these procedures the colour of the clothes were measured using a reflectance spectrometer and expressed as the  $\Delta E$  relative to washed clean white cloth. The results are presented in the table below

10

[Acid red 51] g/L	BC1	White - irradiated	White -dark
0	19.1	1.9	-
0.001	17.5	1.2	2.1
0.002	16.4	0.9	4.2

Clear photo-bleaching of the BC1 stain is observed. A small amount of red colouration is transferred to the white cloth. This red colouration is removed by exposure to sunlight.

15

#### Example 2

A 1g/L SDS surfactant stock solution in water was created. The solution was divided in two and the two halves, and  
20 buffered to pH 10 and pH 8 respectively using standard salts (Hydrion <sup>TM</sup> buffer, purchased from Sigma-Aldrich). The solution was used to wash BC1 stains as follows. A 3.7g piece of BC1 stained cotton cloth plus a 0.7g piece of clean white woven cotton cloth were agitated in 100ml of wash  
25 solution for 20 mins, rinsed twice, then the BC1 cloth irradiated in a WOM for thirty minutes. The  $\Delta E$  of the



- 16 -

clothes relative to a clean standard was then measured. The clothes were then irradiated for a further 30 minutes and remeasured.

- 5 The experiment was repeated using wash solution containing different amounts of Food red 14. The optical density (5cm) of the wash solutions were measured at the lambda (max) of the dye (527nm)
- 10 All the white clothes were tumbled dried in the dark and the pick up of red dye measured by measuring the  $\Delta E$  of the clothes relative to a clean undyed standard.

The results are reported in the table below:

15

ppm dye in solution	pH	Optical density	$\Delta E(BC1)^1$ 30 mins	$\Delta E(BC1)^1$ 60 mins	$\Delta E$ (white cloth) <sup>2</sup>
0	8	0	17.3	16.9	0.5
0.1	8	0.08	16.0	15.7	1.1
0.3	8	0.16	14.8	14.4	2.3
0.8	8	0.36	14.2	13.7	4.7
3.8	8	1.78	10.6	9.9	14.6
0	10	0.0	17.2	16.8	0.5
0.1	10	0.09	15.8	15.4	0.9
0.3	10	0.14	15.0	14.6	1.9
0.8	10	0.36	12.1	11.7	4.1
3.8	10	1.71	9.4	8.8	13.8

Table

<sup>1</sup>average of 4 independent measurements

<sup>2</sup>average of 2 independent measurements

5

Photobleaching of the BC1 stain was observed from all solution containing food red 14. Notably this occurs even at extremely low levels of the dye, 0.1ppm, at this level the dye is barely visible in solution and no red colour could be  
10 discerned by eye on the white cloth. However the BC1 cloth was visibly bleached. The red colour on the white cloth was removed simply by leaving in natural light.

**Example 3**

15

Stains were created on white woven cotton by: placing 1 drop of (a) a saturated solution of turmeric in soya oil or (b) placing 2 drop of Brazilian palm oil.

20 Three of each type of stains (a), (b), BC1 stains and four clean white ballast pieces (total weight =11.4g) were wash at 40°C for 20 minutes in 200 ml of water containing 7g/L of Persil Colour™ washing powder. Persil Colour™ contains less than 5% soap. Polycarboxylate and phosphonate, 5 to 15%  
25 anionic and non-ionic surfactants, 15-30% zeolite and also contains enzymes. Following the wash the cloths were rinsed once in water, once in an acidic solution, and finally once more in water. They were then all irradiated in a WOM for 36 minutes. The residual staining and relative to clean white  
30 cloth measured using a reflectometer and expressed as the  $\Delta E$  value. The discoloration of the white ballast due to

transfer of coloured stain matter in the wash was measured in an analogous manner.

5 The experiment was repeated but with the following, added to the wash solution:

- (a) 1 ppm food red 14
- (b) 10 ppm vitamin K3 (a radical photo-initiator which bleaches carotenoid stains)
- (c) 1 ppm food red 14 plus 10 ppm vitamin K3

10

The results are displayed in the table below

	$\Delta E$			
	BC1	Turmeric	Palm oil	White cotton
Control	16.8	22.5	23.9	2.9
(a)	15.5	22.4	22.8	2.2
(b)	16.6	20.7	8.0	2.5
(c)	15.0	19.3	7.6	1.5

#### Table

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The combination of the Food red 14 with the vitamin K3 gives good overall bleach effects.

#### **Example 4**

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A 3.7g piece of BC1 stained cotton cloth was washed in 80ml of a pH=10 buffered solution of 0.2g/L SDS. The wash consisted in a 15 minute agitation followed by rinsing. The cloth was then irradiated in a WOM for 30 minutes and the

residual staining measured as the  $\Delta E$  relative to clean white cotton. The experiment was also performed with wash liquors containing various xanthene type dyes. The % deposition of the dyes onto the cloth was obtained by measuring the UV-VI spectrum of the wash liquors before and after washing. The results are shown in the table below.

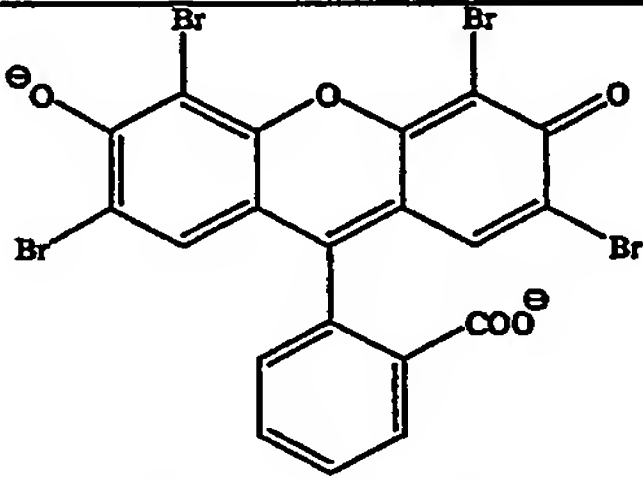
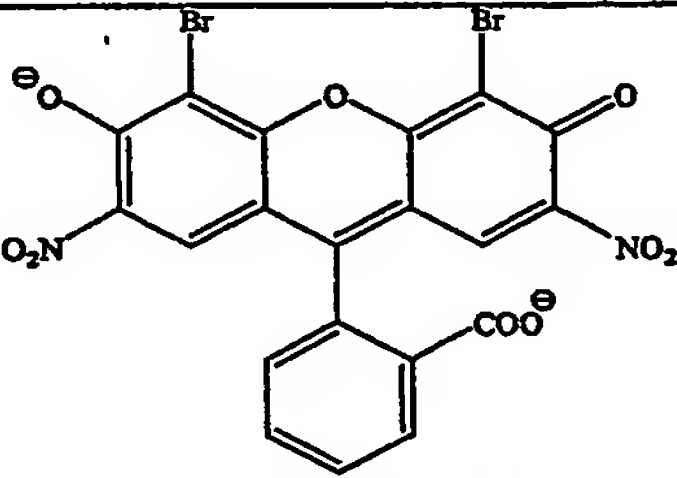
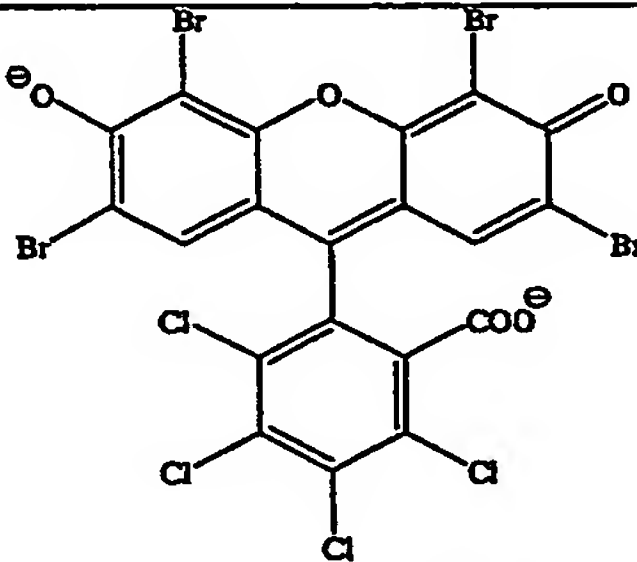
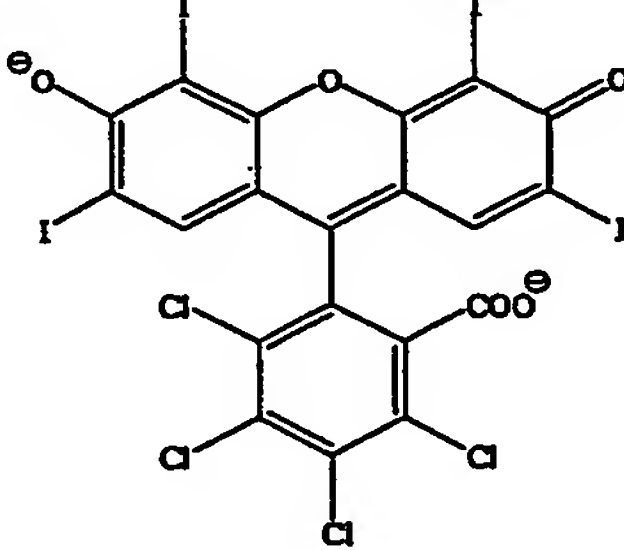
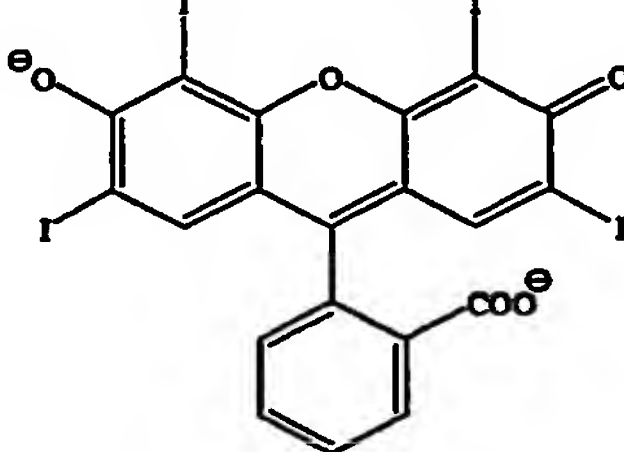
Dye	Amount of dye	BC1 $\Delta E^1$	% deposition
Control	-	17.7	-
Eosin B	2.2ppm	18.1	35
Eosin Y	2.2ppm	12.9	52
Phloxine B	2.1ppm	14.0	9
Rose Bengal	2.2ppm	10.8	19
Food Red 14	2.2ppm	10.9	44

Table

10

<sup>1</sup>average of 4 independent measurements

From the results Food red 14 and Rose Bengal provide superior bleaching.

Name	Dye Anion Structure
Eosin B	 <p>The structure of Eosin B anion consists of a xanthene core. The left benzene ring has a bromine atom at position 6 and a sulfonate group (SO<sub>3</sub><sup>-</sup>) at position 7. The right benzene ring has a bromine atom at position 2 and a carbonyl group (C=O) at position 3. A phenyl ring is attached at position 4, with a sulfonate group (SO<sub>3</sub><sup>-</sup>) at its 2-position.</p>
Eosin Y	 <p>The structure of Eosin Y anion is similar to Eosin B, but the bromine atoms at positions 6 and 2 are replaced by nitro groups (NO<sub>2</sub>). The sulfonate groups (SO<sub>3</sub><sup>-</sup>) are located at positions 7, 3, and on the phenyl ring at its 2-position.</p>
Phloxine B	 <p>The structure of Phloxine B anion features a xanthene core. The left benzene ring has a bromine atom at position 6 and a sulfonate group (SO<sub>3</sub><sup>-</sup>) at position 7. The right benzene ring has a bromine atom at position 2 and a carbonyl group (C=O) at position 3. A 2,4,6-trichlorophenyl ring is attached at position 4, with a sulfonate group (SO<sub>3</sub><sup>-</sup>) at its 1-position.</p>
Rose Bengal	 <p>The structure of Rose Bengal anion is similar to Phloxine B, but the bromine atoms at positions 6 and 2 are replaced by iodine atoms (I). The sulfonate groups (SO<sub>3</sub><sup>-</sup>) are located at positions 7, 3, and on the phenyl ring at its 1-position.</p>
Food Red 14	 <p>The structure of Food Red 14 anion is similar to Rose Bengal, but the iodine atoms at positions 6 and 2 are replaced by bromine atoms (Br). The sulfonate groups (SO<sub>3</sub><sup>-</sup>) are located at positions 7, 3, and on the phenyl ring at its 2-position.</p>



**Example 5**

A stock solution of 1.5g/L of a base washing powder in water was created. The washing powder contained 18% NaLAS, 73% salts (silicate, sodium tri-poly-phosphate, sulphate, carbonate), 3% minors including perborate, fluorescer and enzymes, remainder impurities and water. The solution was divided into 60ml aliquots and various combination of food red 14 dye with acid blue 29 added to this in amount as indicated in the results table. 1 piece of bleached, non-mercerised, non-fluorescent woven cotton cloth (ex Phoenic Calico) weighing 1.3g was placed in the solution at room temperature (20°C). This cloth represents a slightly yellow cotton. The cloth was left to soak for 45 minutes then the solution agitated for 10 mins, rinsed and dried in the dark. After the washes the Ganz whiteness of the cloth was measured (see "assessment of Whiteness and Tint of Fluorescent Substrates with Good Interinstrument Correlation" *Colour Research and Application* 19, 1994). A higher value of Ganz is associated with whiter cloth. The results are the average of 2 experiments.

Food Red 14/ ppm	Acid Blue 29/ ppm	Ganz
0	0	147
0.18	0	150
0.135	0.058	150
0.09	0.117	158
0.045	0.175	162
0	0.233	170
0.09	0	146
0.067	0.029	152
0.045	0.058	156
0.022	0.087	160
0	0.117	163

**Table**

- 5 Mixture of food red 14 with the blue dyes gives an increase in the whiteness of the cloth.

The dye acid black 1 gave similar results.

10 **Example 6**

The experiment of example 5 was repeated. Except here one 3.7g piece of BC1 stained cotton cloth was washed in 60 ml of solution containing various amounts of food red 14 and  
15 acid blue 29. The wash consisted of 30 minutes of agitation. Following the rinses the clothes were irradiated in a WOM for 30 minutes and the residual stain measured as

the  $\Delta E$  relative to clean white cloth. The results are shown in the table below.

Food red 14/ppm	Acid blue 29/ppm	BC1 $\Delta E^1$
0	0	16.9
0.18	0	16.2
0.135	0.058	16.1
0.09	0.117	16.4
0.045	0.175	16.7

5 Table

<sup>1</sup>average of 4 independent measurements.

Hence the mixture of dyes gives photo-bleach effects and  
10 shading benefit (example 5).

**Example 7**

A solution of 0.2g/L SDS surfactant was created in pH=10  
15 buffer (Hydrion <sup>TM</sup>). The solution was divided in 2 and  
RoseBengal added to one half such that its Optical density  
at the maximum absorption of the dye in the visible was 0.81  
(5cm). A similar solution was created for Food Red 14 but  
with a higher optical density of 0.93. The Solution of Food  
20 Red 14 under indoor lighting was much paler to the eye than  
that of Rose Bengal. This was confirmed by 12 independent  
adults aged in the range 25-60 years. All stated the Rose  
Bengal solution to have a more intense colour, despite the  
fact it had a lower optical density. This clearly shows the

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advantage in being able to dose Food red 14 at a higher level without over colouring the liquor.

## We claim:

1. A bleaching composition comprising:
  - a) from 0.0001 wt/wt% to 0.1 wt/wt% of Food red 14 dye;
  - 5 b) from 0 to 40% other bleaching species; and,
  - c) the balance carriers and adjunct ingredients to 100 wt/wt % of the total bleaching composition, wherein upon addition of a unit dose of the bleaching composition to an aqueous environment a wash liquor having a pH in the
  - 10 range 8 to 11 is provided.
2. A bleaching composition according to claim 1, wherein the Food red 14 bleaching composition is present in the range of 0.001 to 0.01 wt/wt%.
- 15 3. A bleaching composition according to claim 1 or 2, wherein the bleaching composition comprises other photo-bleaches.
- 20 4. A bleaching composition according to claim 3, wherein the bleaching composition comprises vitamin K3.
- 25 5. A bleaching composition according to any preceding claim, wherein the bleaching composition comprises an air bleach catalyst or ligand thereof, the bleaching composition comprising less than 1 % wt/wt total concentration peroxy-based or peroxy-generating bleach system.



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6. A bleaching composition according to any preceding claim, wherein the bleaching composition is a liquid bleaching composition.
- 5 7. A bleaching composition according to any one of claims 1 to 4, wherein the bleaching composition comprises a peroxy species.
- 10 8. A bleaching composition according to claim 7, wherein the bleaching composition comprises a bleach activator.
- 15 9. A bleaching composition according to any preceding claim, wherein the bleaching composition comprises from 5 to 40 wt/wt % of a surfactant having an HLB greater than 15.

Abstract of the Invention

The present invention concerns photo-bleaching with a dye and in particular the bleaching of laundry items.

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